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# EVALUATION OF UNION CARBIDE CORPORATION EPOXY BINDER FOR USE IN AIRCRAFT PARACHUTE FLARES



PREPARED BY

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#### NAVAL AMMUNITION DEPOT Crane, Indiana 47522

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#### EVALUATION OF UNION CARBIDE CORPORATION EPOXY BINDER FOR USE IN AIRCRAFT PARACHUTE FLARES

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#### ABSTRACT

The material used as a binder in the pyrotechnic compositions (candles) of Mk 24 and Mk 45 Aircraft Parachute Flares, since the 1968 conversion from Laminac to epoxy, has been procured on a solesource basis from the Dow Chemical Company. A thorough analysis of all pertinent properties of an epoxy produced by Union Carbide Corporation has demonstrated that this material is, in all significant respects, as good as that obtained from Dow. This report recommends that Union Carbide Corporation be approved as an alternate supplier for the epoxy used in these flare candles.

#### TABLE OF CONTENTS

																						Ī	age	
Backgr	round	٠.			•			•										•			•	•	1	
Chemic				on	of	FI	Jn:	ior	1 (	Car	rbi	d	e I	Ерс	oxy	7.							1	
Re	esin.																						1	
На	rdene	r.			•	•	•			•			•	•	•	٠	•	•	•	•	•	•	2	
Items	Evalu	ate	1					•			•			•	•								2	
Cand1e	Form	ulat	ior	ıs.								•								•			3	
Gas Ge	enerat	ion	Tes	st.																			3	
Compat	ibili	ty !	act	ors	5.						4												4	
Carlo H		7																						
Physic	al Pr	oper	rtie	es.																			4	
Mi	grati	on ?	rest	s.																			5	
Th	nermal	Deg	grad	lati	ior	1 7	re:	sts	5.	•	•		•		•		•	•	•	•	•		6	
Conclu	sions					•													•	•		•	6	
Recomm	nendat	ion						•						•						•			7	
TABLE	I - 0	and	le I	orr	nu]	at	tic	ons	5 (	Cor	nta	ii	niı	ng	Ur	nic	on	Ca	art	oio	le			
		poxy													•			•	•	•		•	8	
TABLE	II.,-,	Stat	cic	Fui	nct	ii	n	ing	3	Гes	st				•								9	
TABLE	TTT	C+.	+ :	т.	201		D	· f			no	1	1 5	- 0	n 1	rei	1							
TABLE	111 -		ndit																				11	
TABLE	IV -	Pres	ssui	re l	Bui	110	l-l	Jp	Te	est	· .												13	
TABLE	V - 1	ens	ile	St	rei	ngt	th	Te	es	t.													14	
TABLE	VI -	The	rmal	l De	egi	rac	lat	tic	on	Te	est												15	
Thermo	grams																					10	6-35	

#### EVALUATION OF UNION CARBIDE CORPORATION EPOXY BINDER FOR USE IN AIRCRAFT PARACHUTE FLARES

#### 1. Background.

- a. This study was undertaken for the express purpose of evaluating the epoxy binder material produced by and available from the Union Carbide Corporation. Since 1968, the Dow Chemical Company has been the designated sole supplier of epoxy binder used in production of illuminating candle composition for Mk 24 Mod 4 and Mk 45 Mod 0 Aircraft Parachute Flares. There is an obvious advantage to the Government in having two or more suppliers on a competitive basis.
- b. The Union Carbide Corporation supplied the binder for the evaluation tests reported herewith and quoted a price for large-quantity procurement which is significantly lower than the price being paid for the presently used epoxy.
- 2. Chemical Description of Union Carbide Epoxy. The epoxy binder material produced by Union Carbide Corporation consists of a two-part system—resin and hardener. These two prime constituents are described as follows:
- a. Resin. The resin, which is identified as ERLA-2713, is a diepoxide which has been modified for low viscosity. It is a reactive diluent, general purpose-type liquid. The reactive diluent, approximately 20% by weight, is CGE (cresylglycidye ether), and the resin is based upon the reaction of bisphenol-A and epichlorohydrin to produce diglycidyl ether of bisphenol-A as the major constituent. The epoxy equivalent weight range of 185-195 and low viscosity of 500-700 cps is practically the same as used in the present production binder resin,

Dow's DER-321.

b. <u>Hardener</u>. The curing agent or hardener, identified as ZZLA-0672, is a moderately low viscosity, aliphalic polyamine with an amine hydrogen equivalent weight of 104.5. The epoxy resin is cured with this reactive crosslinking agent TETA(triethylene tetramine) which has been modified by the addition of a nonreactive diluent, mobilsol-44, for improved flexibility. This amine hardener is a room-temperature-curing type which makes the application of heat unnecessary; however, a short post cure at high temperature (100°C) will improve the properties of the cured binder. To obtain optimum properties and to insure complete cure of the resin, it is desirable to react the resin and curing agent at approximate stoichiometric quantities. To determine the ratio of resin to hardener, calculations were made as follows:

\*Phr of hardener (0672) = 
$$\frac{\text{Amine H eq. wt. X } 100}{\text{Epoxy eq.wt.}}$$

Phr (0672) - 
$$\frac{(104.5)(100)}{190}$$
 = 55.0

Therefore the mixing ratio which is used for this binder system is 100 Phr of ERL-2713 to 55 Phr of ZZLA-0672.

- 3. <u>Items evaluated</u>. The following items and materials were tested during this evaluation:
- a. Complete flare candles made for candlepower, burn time, temperature and humidity, vibration, and pressure buildup tests.
- b. Separate binder constituents (resin and hardener) for vapor pressure and compatibility tests.
- c. Cured Dow and Union Carbide epoxy for physical properties determination and migration tests.

- 4. Candle formulations. Table I gives the formulations of candle compositions containing both 3.5% and 4% binder material. The six separate formulas resulted from varying the magnesium content of the batches. Table II shows the results of burn time and candlepower tests that were conducted using candles that were made from these six formulas as well as those made from the standard Mk 24 and Mk 45 formulas. is readily seen from Table II that candles made with Union Carbide and standard, that is, presently used epoxy binders, are virtually equal from the standpoint of candlepower. The relative intensity of the light energy output varies in direct proportion with the percentage of magnesium in the formula. Thus, the total formulation has to be taken into account in comparing the test data. Formula "C" has the same magnesium content as the control candles. In T&H testing, it was noted that the loss in burn time, which is characteristic of this type of testing, was significantly less in the Union Carbide candles than it was in the control candles. Table III shows that the Union Carbide candles had a variation of only 3 to 5 seconds, whereas the burning time of the control candles varied 10 to 12 seconds after T&H cycling.
- 5. <u>Gas Generation Test.</u> A pressure buildup-type test was conducted by placing the test candles in sealed tubes with pressure gages attached to their tops. These tubes were then placed in a temperature-controlled chamber which was set at 140°F. The gages were read and the data recorded each day for 10 days with results as shown in Table IV. This data indicates that the Union Carbide candles generated an average pressure of 3.25 psi as compared with 3.125 psi average for the control candles. The primary significance here is not in the comparison of one

epoxy with another; however, it is indicative of the desirability of epoxy from whatever source. This relatively low generation of hydrogen is an excellent property of the epoxies - they provide good protection which effectively prevents absorption of water into the flare composition. 6. Compatibility Factors. The separate chemical constituents of the Union Carbide epoxy; that is, the resin and the hardener, proved to be totally compatible with other flare components with which they might come into contact. There developed, however, a cause for some concern in respect to the nonreactive diluent, Mobilsol-44, which is simply a long-chained hydrocarbon compound that is added to the Union Carbide hardener to increase flexibility. Since this ingredient does not react in the fully cured system, it was feared that there might be a possibility of its separating from the binder through evaporation at higher temperatures. A vapor-pressure test was conducted on the diluent to investigate this possibility. The pressure was determined over a temperature range of from 150°F to 324.5°F as shown below. (No measurable pressure was obtained at temperatures below 150°F). A vapor pressure buildup of 6.6 mm of mercury at 190°F is considered to be acceptably low for illuminating

Temperature	°F	Vapor	Pressure,	Torr
150			3.5	
190			6.6	
260			17.0	
324.5			26.0	

flare applications.

7. Physical Properties. The physical properties that were investigated showed a wide variation between the two binder systems. Table V clearly demonstrates the very pronounced differences in elasticity. The Union Carbide epoxy is rubbery and quite flexible, whereas the epoxy obtained

from the Dow source is comparatively rigid and brittle. This, of course, accounts for the rather extreme differences in tensile strength and stretching properties as shown in Table V. This elasticity in the Union Carbide product results from the inclusion in its hardener formulation of the nonreactive diluent. Mobilsol-44 was added for the express purpose of achieving the flexibility that was noted. Such flexibility, however, is gained at the expense of tensile and impact strength. The chief requisites for a nonreactive diluent are (1) that it not foam or vaporize during the curing process, and (2) that it not migrate from the fully cured composition.

a. Migration Tests. The Quality Evaluation Department at Crane conducted migration tests on the cured Union Carbide binder material. In these tests, an attempt was made to measure the rate of migration of volatiles from the cured binder at a temperature of 158°F. This test would show if any of the nonreactive diluent, which is not tied up in the cured resin structure, comes out of the cured system. The test consisted of using a 1" block of cured resin against a 1" block of steel with a porous asbestos paper disc sandwiched in between. The asbestos discs were carefully weighed and on some control units a drop of Mobilsol-44 was added. During the tests the control samples lost their charge of Mobilsol-44 and when the asbestos discs were reweighed, no difference in weight could be detected. Therefore, it could not be determined if any of the nonreactive diluent came out of the system and evaporated, or remained intact. Since thermal degradation tests were in progress, it was decided not to rerun the migration tests as these tests would show the loss of weight of the cured resin which would indicate

any evaporation of the nonreactive diluent.

- b. Thermal Degradation Tests. Table VI gives detailed results of thermal degradation tests which were conducted on both the Union Carbide epoxy and the Dow epoxy presently used in flares. It was evident that these tests would show any migration of the nonreactive diluent out of the Union Carbide system and thus supply the information that was unobtainable through the migration tests. Cured binders were tested at 75°C, 100°C, and 160°C. As Table VI shows, the diluent exuded at 75°C (167°F) with a weight loss of approximately 2%. This increases as the temperature is raised to 160°C. Since the diluent constitutes 30 to 33 percent of the Union Carbide binder weight, a loss of the magnitude of 2% at 167°F is considered to be not harmful. The weight loss in the control epoxy system was .5 to 1.0 percent at 75°C which indicates that a nonreactive diluent is present in this system also. Neither of the epoxies foamed during the curing process.
- 8. <u>Conclusions</u>. A binder's main purpose in illuminating flare compositions is to provide a bond for holding the fuel and oxidizing constituents of the candle together. It should also have the desirable properties of insulating against moisture absorption, of increasing the quantity of mix per volumetric unit, of regulating to a degree the candle's burning time, and of acting as a wetting agent to desensitize the magnesium during the mixing process. The Union Carbide epoxy binder amply satisfies all of these requirements. While it proved to be physically weaker than the presently used epoxy, its strength is more than adequate for the intended purpose since candles are consolidated under high pressure into heavy cardboard tubes which aid greatly in holding the ingredients together.

The greater flexibility of the Union Carbide product should, it is believed, add to its value as a binder by reducing the tendency of candle composition to "chunk out" during burning. The Union Carbide epoxy showed improvement over the presently used binder in that it exhibited a more controllable burning rate. It is also readily available at a lower cost.

9. Recommendation. It is recommended that the Union Carbide Corporation be approved as an alternate supplier of binder epoxy for the Mk 24 and Mk 45 Aircraft Parachute Flare production and that the drawings for both of these flares be amended to include the Union Carbide epoxy as an alternate binder material.

Candle Formulations Containing Union Carbide Epoxy

			Formulas	as		
	А	В	U	D	ш	ш
Used for Test Candles Nos.	1-36	40-46	99-09	57-79	98-08	96-06
Magnesium Weight (in pounds)	70	71,25	72.5	70	71,25	72.5
Magnesium Percentage	29%	21%	28%	%95	21%	28%
Oxidizer Weight (in pounds)	50,625	49,375	48,125	20	48,75	47.5
Oxidizer Percentage	40.5%	39.5%	38.5%	40%	39%	38%
Binder Percentage	3.5%	3.5%	3.5%	4%	4%	4%
Binder Resin Weight (in grams)	1,281	1,281	1,281	1,460	1,460	1,460
Binder Hardener Weight (in grams)	705	705	202	802	802	805

For purpose of these tests (see Table II), Control candles were standard Mk 24 and Mk 45 components. they were numbered 100 and above. NOTE:

TABLE II Static Functioning Test

11	FORMULA A CAND	LES		FORMULA B CANDI	LES
Candle No.	Candlepower	Burn Time (sec.)	Candle No.	Candlepower	Burn Time (sec.)
12	1,837,500	204.0	44	1,819,500	208.0
23	1,791,000	204.0	43	1,845,500	209.0
28	1,636,000	206.0	40	1,681,500	207.0
6	1,789,500	210.0	41	1,824,000	209.0
15	1,884,500	209.0	46	1,870,500	210.0
Average	1,787,700	206.6		1,808,200	208.6
Control Candle Avg.	1,865,900	204.2		1,865,900	204.2

	FORMULA C CANDI	LES	<u> </u>	ORMULA D CANDI	D CANDLES		
Candle No.	Candlepower	Burn Time (sec.)	Candle No.	Candlepower	Burn Time (sec.)		
53	1,884,000	203.8	68	1,760,500	223.0		
51	1,820,500	205.5	74	1,726,000	225.0		
56	1,798,000	202.0	57	1,600,000	225.0		
52	1,859,000	205.0	63	1,698,000	221.0		
54	1,969,500	205.0	67	1,827,000	225.0		
Average	1,866,200	204.0		1,722,300	223.8		
Control Candle Avg.	1,865,900	204.2		1,865,900	204.2		

## TABLE II (cont.) Static Functioning Test

	FORMULA E CANDI	LES		FORMULA F CANDI	LES
Candle No.	Candlepower	Burn Time (sec.)	Candle No.	Candlepower	Burn Time (sec.)
86	1,920,500	193.0	90	1,914,500	204.0
81	1,934,500	198.0	93	1,859,000	210.5
80	1,784,000	195.0	92	1,666,500	218.0
85	1,856,500	202.0	94	1,807,000	211.0
82	2,093,000	200.0	95	1,894,500	212.0
Average	1,917,700	197.6		1,828,300	211.1
Control Candle Avg.	1,865,900	204.2		1,865,900	204.2

#### CONTROL CANDLES

Candle No.	Candlepower	Burn Time (sec.)
108	1,878,000	205.0
109	1,824,000	206.0
110	1,840,000	194.0
111	1,898,000	202.0
112	1,889,500	214.0
Average	1,865,900	204.2

TABLE III
Static Tests Before And After T&H Conditioning

#### FORMULA A CANDLES

No	t Conditione	<u>:d</u>	<u>A</u>	fter 14-Day T	<u>&amp;Н</u>			
Candle No.	Candlepower	Burn Time	Candle No.	Candlepower	Burn Time	Variat	ion	
2	1,896,000	210.0	8	1,853,000	205.0	-5 s	ec.	
21	1,825,500	208.0	32	1,849,500	202.0	-6 s	ec.	
30	1,867,500	204.0	16	1,917,500	204.0	-0 s	ec.	
35	1,817,500	210.0	27	1,886,500	203.0	-7 s	ec.	
7	1,805,000	209.0	1	1,850,000	210.8	+1.8	sec.	
17	1,899,000	207.0	20	1,883,500	205.0	-2 s	ec.	
Averages	1,851,750	208.0		1,873,333	204.977	-3 s	ec.	

#### FORMULA D CANDLES

No	ot Conditione	<u>:d</u>	P	fter 14-Day T	<u>&amp;H</u>	
Candle No.	Candlepower	Burn Time	Candle No.	Candlepower	Burn Time	Variation
62	1,749,500	219.0	75	1,716,500	217.0	-2 sec.
73	1,696,500	226.0	72	1,837,000	222.0	-4 sec.
60	1,707,000	224.0	59	1,704,000	212.6	-11.4 sec.
70	1,802,500	225.0	65	1,717,500	222.0	-3 sec.
Averages	1,738,875	223.5		1,743,750	218.4	-5.1 sec.

### TABLE III (cont.) Static Tests Before And After T&H Conditioning

#### CONTROL CANDLES

No	ot Conditione	d				
Candle No.	Candlepower	Burn Time	Candle No.	Candlepower	Burn Time	Variation
102	2,151,500	199.0	100	2,075,000	182.0	-17 sec.
103	2,020,500	201.0	101	2,044,500	193.0	-8 sec.
Averages	2,086,000	200.0		2,059,750	187.5	-12.5 sec.

TABLE IV Pressure Build-Up Test

#### Union Carbide Epoxy Candles

#### Daily Pressure Measurements (psi)

Ca	ndle	4	9.6			75	10 mg 2	1			
	No.	lst	2nd	3rd	4th	5th	6th	7th	8th	9th	10th
	4	0	0	0	0	0	0	0	0	1	3
	10	0	0	0	0	. 0	0	0	0	1	2
	25	5	4	4	3	4	4	3	5	5	6
	31	2	1	0	0	1	1	1	2	3	4
	42	4	2	1	1	2	2	2	3	4	4.5
	45	3	1	1	0	0	0	0	1	2	4
	50	0	0	Ú	0	0	0	0	0	1	3
	55	0	0	0	0	0	0	0	0	0	0
	64	3	2	1	0	1	1	+1	2	2	4
	69	0	0	0	0	0	0	0	0	0	1
	83	3	2	2	1	3	3	3	5	6	7
	84	2	0	0	0	0	0	0	0	1	3
	91	0	0	0	0	0	0	0	1	1	3
	96	0	0	0	0	0	0	0	0	0	1

Average 10th Day Pressure 3.25

#### Standard Mk 45 Candles

#### Daily Pressure Measurements (psi)

104	1	0	0	0	0	0	.5	2	3	4	
105	0	0	0	0	0	0	0	0	0	1	
106										3	
107	3	2	1	0	1	1	1	2	3	4.5	

Average 10th Day Pressure 3.125

#### TABLE V Tensile Strength Test

Epoxy	Breaks at (psi)	Elongation (%)
Dow	3300	11
Union Carbide	610	60

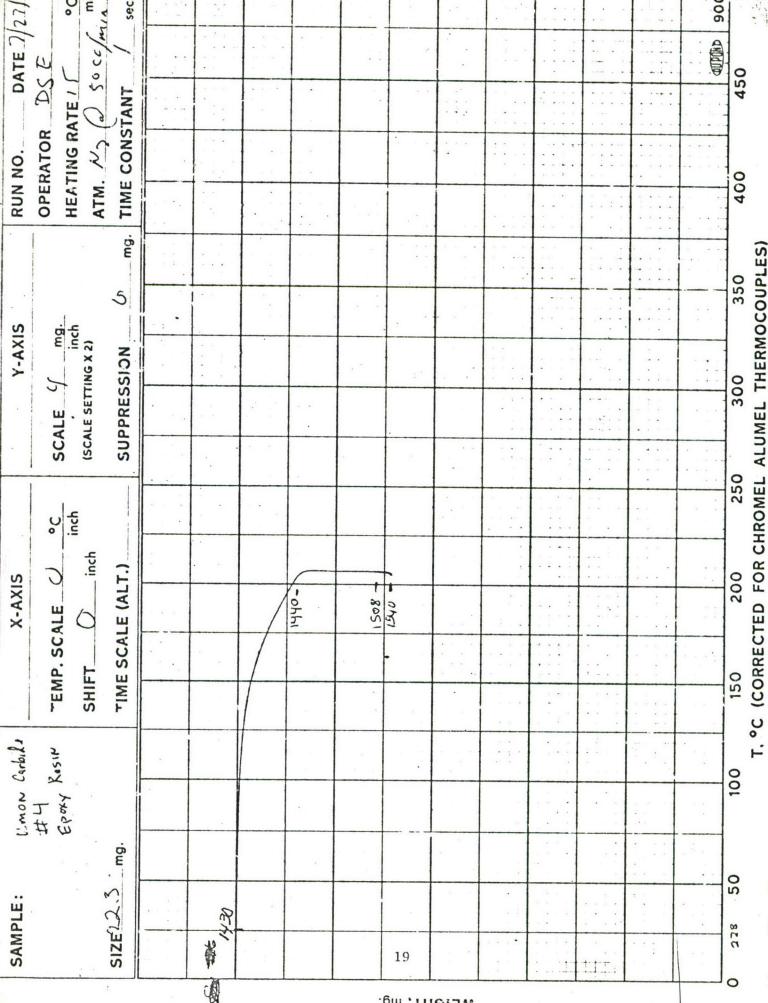
TABLE VI Thermal Degradation Test

Epoxy Sample	
Control (Dow) 2 .5 - 1.0 .5 - 1.0 2.0 - Control (Dow) 3 .5 - 1.0 .5 - 1.0 2.0 - Control (Dow) 4 .5 - 1.0 .5 - 1.0 2.0 - Control (Dow) 5 .5 - 1.0 .5 - 1.0 2.0 - Control (Dow) 6 1.0 - 2.0 1.0 - 2.0 10.0 - Control (Dow) 7 1.0 - 2.0 1.0 - 2.0 15.0 - Control (Dow) 8 .5 - 1.0 1.0 - 2.0 10.0 - Control (Dow) 9 .5 - 1.0 1.0 - 2.0 10.0 - Control (Dow) 9 .5 - 1.0 1.0 - 2.0 10.0 - Control (Dow) 10 1.0 - 2.0 1.0 - 2.0 10.0 - Control (Dow) 10 2.0 - 10.0 15.0 - 20.0 Union Carbide 1 2.0 - 10.0 15.0 - 20.0 Union Carbide 3 2.0 - 10.0 10.0 - 15.0 Union Carbide 5 2.0 - 10.0 10.0 - 15.0 Union Carbide 6 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0	
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Control (Dow) 7 1.0 - 2.0 1.0 - 2.0 15.0 - Control (Dow) 8 .5 - 1.0 1.0 - 2.0 10.0 - Control (Dow) 9 .5 - 1.0 1.0 - 2.0 2.0 - Control (Dow) 10 1.0 - 2.0 1.0 - 2.0 10.0 - Union Carbide 1 2.0 - 10.0 15.0 - 20.0 Union Carbide 2 2.0 - 10.0 20.0 - 20.0 Union Carbide 3 2.0 - 10.0 10.0 - 15.0 Union Carbide 4 2.0 - 10.0 10.0 - 15.0 Union Carbide 5 2.0 - 10.0 10.0 - 15.0 Union Carbide 6 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0	10.0
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Control (Dow)       9       .5 - 1.0       1.0 - 2.0       2.0 -         Control (Dow)       10       1.0 - 2.0       1.0 - 2.0       10.0 -         Union Carbide       1       2.0 - 10.0       15.0 - 20.0         Union Carbide       2       2.0 - 10.0       20.0 - 20.0         Union Carbide       3       2.0 - 10.0       10.0 - 15.0         Union Carbide       4       2.0 - 10.0       10.0 - 15.0         Union Carbide       5       2.0 - 10.0       10.0 - 15.0         Union Carbide       6       2.0 - 10.0       10.0 - 15.0         Union Carbide       7       2.0 - 10.0       10.0 - 15.0	20.0
Control (Dow)       10       1.0 - 2.0       1.0 - 2.0       10.0 - 2.0         Union Carbide       1       2.0 - 10.0       15.0 - 20.0         Union Carbide       2       2.0 - 10.0       20.0 - 20.0         Union Carbide       3       2.0 - 10.0       10.0 - 15.0         Union Carbide       4       2.0 - 10.0       10.0 - 15.0         Union Carbide       5       2.0 - 10.0       10.0 - 15.0         Union Carbide       6       2.0 - 10.0       10.0 - 15.0         Union Carbide       7       2.0 - 10.0       10.0 - 15.0	15.0
Union Carbide 1 2.0 - 10.0 15.0 - 20.0  Union Carbide 2 2.0 - 10.0 20.0 - 20.0  Union Carbide 3 2.0 - 10.0 10.0 - 15.0  Union Carbide 4 2.0 - 10.0 10.0 - 15.0  Union Carbide 5 2.0 - 10.0 10.0 - 15.0  Union Carbide 6 2.0 - 10.0 10.0 - 15.0  Union Carbide 7 2.0 - 10.0 10.0 - 15.0	10.0
Union Carbide 2 2.0 - 10.0 20.0 - 20.0  Union Carbide 3 2.0 - 10.0 10.0 - 15.0  Union Carbide 4 2.0 - 10.0 10.0 - 15.0  Union Carbide 5 2.0 - 10.0 10.0 - 15.0  Union Carbide 6 2.0 - 10.0 10.0 - 15.0  Union Carbide 7 2.0 - 10.0 10.0 - 15.0	15.0
Union Carbide 3 2.0 - 10.0 10.0 - 15.0  Union Carbide 4 2.0 - 10.0 10.0 - 15.0  Union Carbide 5 2.0 - 10.0 10.0 - 15.0  Union Carbide 6 2.0 - 10.0 10.0 - 15.0  Union Carbide 7 2.0 - 10.0 10.0 - 15.0	20.0
Union Carbide 4 2.0 - 10.0 10.0 - 15.0 Union Carbide 5 2.0 - 10.0 10.0 - 15.0 Union Carbide 6 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0	20.0
Union Carbide 5 2.0 - 10.0 10.0 - 15.0 Union Carbide 6 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0	20.0
Union Carbide 6 2.0 - 10.0 10.0 - 15.0 Union Carbide 7 2.0 - 10.0 10.0 - 15.0	20.0
Union Carbide 7 2.0 - 10.0 10.0 - 15.0	20.0
	20.0
200 250	20.0
Union Carbide 8 2.0 - 10.0 10.0 - 15.0	20.0
Union Carbide 9 .5 - 1.0 2.0 - 10.0	20.0
Union Carbide 10 2.0 - 10.0 2.0 - 10.0	20.0

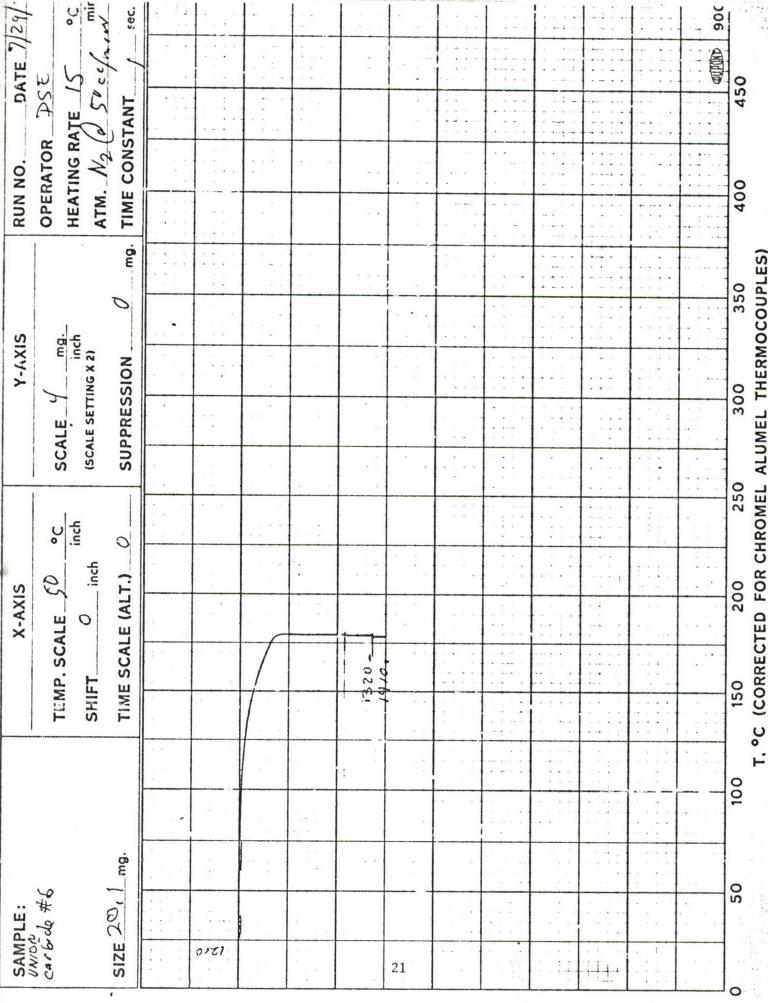
	SAME	LE: L	ory Re	NIS.				X-AXIS	ıs				Y-AXIS	S		RUN NO.	NO.	D	FE	15/2
	Union	Union Carbide Botch 1	Bet	4	-0	TEM	TEMP. SCALE	1	50	O inch	SC/	SCALE H mg.	E . É	æ-5		OPE	OPERATOR HEATING RATE	M	5	00
,						SHIFT	-	0	- inch		(SCA	ILE SETT	ING X 2)			ATM.		N2 @ 504,	4/min	3
	SIZE 216	216	-mg.			TIME	TIME SCALE (AI	E (AL	LT.)		SU	SUPPRESSION	SION	P	mg.		TIME CONSTANT	TANT	-	sec
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0		20	0	5	100 T.	150 T, °C (CORRECTED	O	N	200 250 FOR CHROMEL	2 IROME	250 MEL ALL	3( JMEL	300 L THERN	34000	300 350 ALUMEL THERMOCOUPLES)	400	0	4	450	1.16

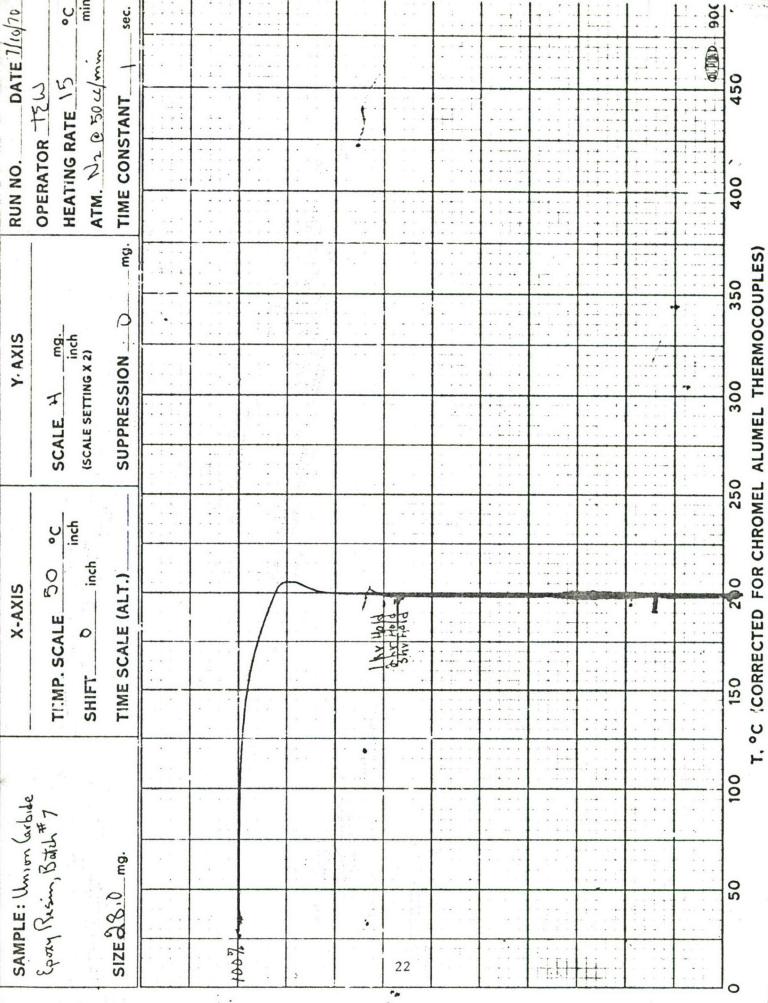
	SAME	SAMPLE: E P	E PUXY RESIN BIX #>	>		X-AXIS	XIS				Y-AXIS	(IS		RUI	RUN NO.		DATE 7	123/
7,		,			TEMP. SCALE	CALE	50	oCinch	50	SCALE 4	SCALE 4 mg.	ng.		OP HE	OPERATOR DSE HEATING RATE 15	A	157	0.
	SIZE	SIZE 24.4 mg.	.g.	, Par	TIME SCALE (ALT.)	ALE (A	LT.)		ns	IPPRE	SUPPRESSION	0	٤	ATA mg. TIM	ATM. N (2)	3 A	50 05	m, m
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)		Oc	T 100	T, °C (	150 (CORRECTED	C	200 FORCI	250 250 FOR CHROMEL		300 IMEL TH	300 350 ALUMEL THERMOCOUPLES	350 OCOUPI	50 PLES	400	00	4	450	

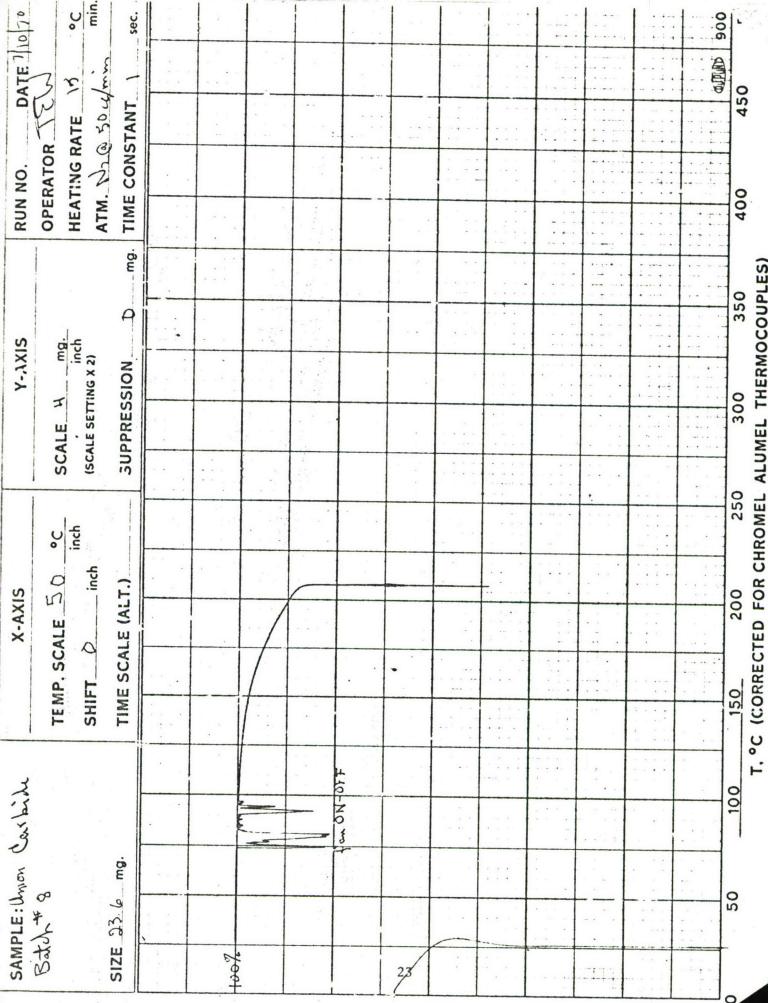
D	SAMPLE:	PLE:		( N 1 C. N		٤		X-AXIS	10				Y-AXIS	S		RUN NO.	NO.	D.	DATE 7 24	24
		c.	•	Carbide #3	e #3	F	FMP SCALE		20	ပို	SCA	SCALE	ű.			OPER	OPERATOR	50	E	
		•		Resin			; ; ;			inch	(SCA	LE SETT	inch (SCALE SETTING X 2)	ج		HE/.T	ING R			0 E
	,		٠			SHIFT			- inch							ATM. Ng	N	(8)	30 cc/m	3.8
	SIZE	70.7	mg.			IME	TIME SCALE (ALT.)	E (AL	(.)		SU	SUPPRESSION	SION	0	.gm		TIME CONSTANT	TANT	-	sec
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)	0	S	50	100		150 C (COR	150 T, °C (CORRECTED	N	OR CF	250 IROMEL	SO L	300 JMEL TH	FOR CHROMEL ALUMEL THERMOCOUPLES	350	O SLES)	400	0	450	0:	11



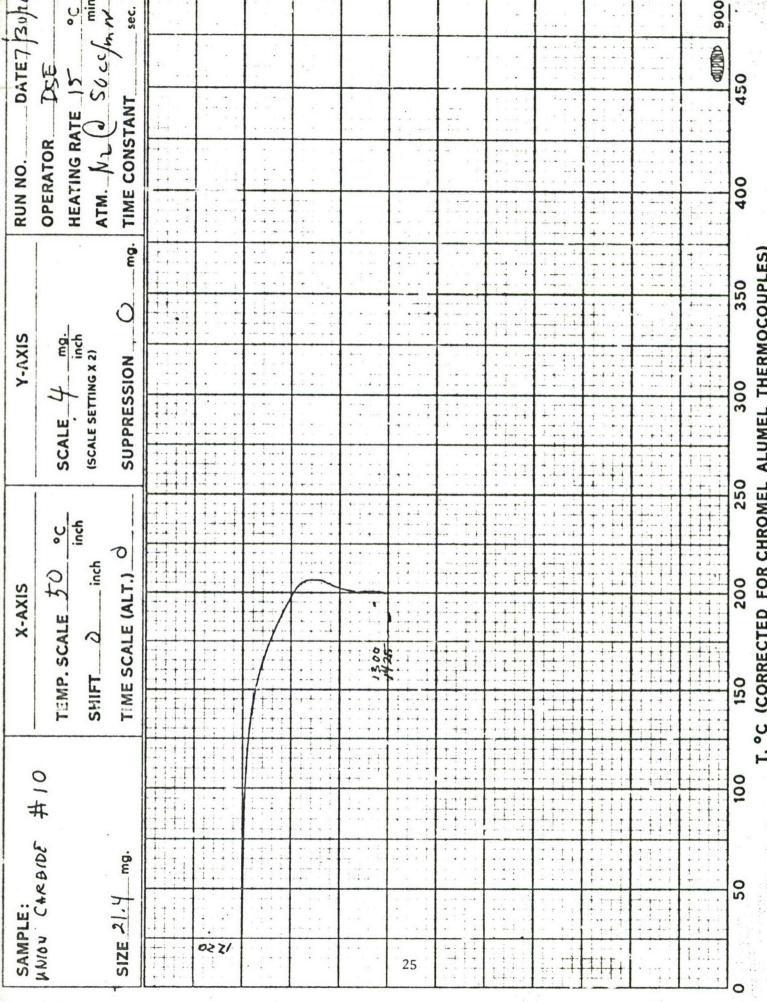
	SAM	PLE:	SAMPLE: Eposy Risin	· *				X-AXI	S				Y-AXIS	IS		RUN NO.	NO.	D	DATE 7 15 70	13/7
	Mura	n Carl	ide 182	tc 43		7.EM	TEMP. SCALE		20	o°C	SC	ALE X	اع			OPE	OPERATOR TELL	127	31	
						SHIFT	l <del>.</del>		100	inch	(SC	inch (SCALE SETTING X 2)	in (S X 2)	ch		HEA	HEATING RATE	ATE	D .	O.E
,		010				5	•									ATM	ATM. No @ 50 4/m	2 50 6	stomes	
	SIZE	SIZE ALL	mg.			TIMIT	E SCA	TIME SCALE (ALT.)	T.)		SU-	SUPPRESSION	SSION	0	mg.		TIME CONSTANT	TANT	7	sec.
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- 1									-											
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9	•	3,	20	10	100 T.°	150 °C (COR	150 (CORRECTED	101	O OR CI	00 250 FOR CHROMEL		3( JMEL	300 L THERN	3,10001	300 350 ALUMEL THERMOCOUPLES)	400	0	4	450	







SAN	SAMPLE:	,	=				X-AXIS	S				Y-AXIS	S		RUN NO.		DATE 7/36/-	1/34
ਤ 	NON Ce	Union Carbida	#		TEM	TEMP. SCALE	1	5.0	0 40.5	SCA	SCALE 4	SCALE 4 mg.	9		OPERATOR HEATING RA	<b> </b>	DSE EVS	0.
		NI NI		٠	SHIFT	-T-		inch		(SCA	LE SETT	ING X 2)	Ę	4	ATM.	ATM. N2 () COCC	/	min
SIZI	SIZE 24.8	mg.			T.M.	TIME SCALE (ALT.)	E (AL		0	SUB	SUPPRESSION	SION	0	.ge		TIME CONSTANT	,	sec.
				11.								:						
2559							:											
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	7							1005.										
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0	5	50	100	1	150 C (COR	150 T. °C (CORRECTED	N	ORC	100 250 300 350 350 FOR CHROMEL ALUMEL THERMOCOUPLES)	SO L ALU	300 IMEL TH	O THERM	350 OCOUPL	O PLES)	400		450	



-	SAME	PLE:			The state of the s		A variable of the constraints	X-AXIS	IS			-	Y-AXIS	S		RUN	RUN NO.	D/	DATE 1/284	1284
	4 4	188 # 188 #	Dow #1 Fronky Resin	2		TEMP.	IP. SC FT	TEMP. SCALE S SHIFT	O in C	°C inch	SCA (SCA	SCALE U mg.	inc NG X 2)	عاد		OPE HEA	OPERATOR DSE HEATING RATE $ S $	ATE S	1	O. mim
•	SIZE	SIZE 3/.8	mg.			MIT	E SCA	TIME SCALE (ALT.)	1	0	SUF	SUPPRESSION	NOIS	0	mg.		TIME CONSTANT	TANT	-	sec.
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0		S	50	100		150 Z	150 CORREC	2 CTED	200 FOR C	300 350 300 350 FOR CHROMEL ALUMEL THERMOCOUPLES)	250 MEL ALU	30C JMEL TH	CHERM	350 OCOUPI	50 PLES)	4	400	4	450	

SAMPLE:	Dow #	#7			X-AXIS					Y-AXIS	. 51	·	RUN NO.	NO.		DATE	1/2
٠.	4		TEM	TEMP. SCALE	LE SO	1	o inch	SC/	SCALE H	E .c	mg.		OPE HEA	OPERATOR HEATING RA	OPERATOR DAE	S	0
			SHIFT	FT		ج		(SCA	ILE SETT	(SCALE SETTING X 2)			ATM	N	ATM. N 2 @ 50 cc.	0 00	Linn
- 11	-mg.		TIM	TIME SCALE (ALT.)	E (ALT			SU	SUPPRESSION	SION	0	mg.		TIME CONSTANT	TANT		se
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_	50	100 T.	15 T, °C (C(	150 (CORPECTED	2	O OR CH	250 IROMEL	SO L	3C	300 L THERN	36	00 250 300 350 FOR CHROMEL ALUMEL THERMOCOUPLES)		400	4	450	

	SAMPLE:						X-AXIS	S				Y-AXIS	IS		RUN	RUN NO.		DATE 1/2/	1751
7	Epoxy Rosin	ROSIN			CEMP.	FEMP. SCALE	O O	0	°C inch	SC	SCALE X	SCALE SETTING X 2)	mg. inch 2)		OPE	OPERATOR HEATING RATE	E (	DSE 18	0
•	SIZE 20,6	€ mg.			TIM	TIME SCALE (ALT.)	E (AL	2		SU	SUPPRESSION	SION	0	.mg.		ATM. N2 (6) TIME CONSTANT	STANT	79 00	1
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0	4	50	7	100 T.°	150 T, °C (CORRECTED	ORREC	10	O OR CH	00 250 FOR CHROMEL	O L ALU	30C	CHERM	30C 350 ALUMEL THERMOCOUPLES)	0 PLES)	400	0	4	450	1

28	, E	m, A.											)6	
DATE 1/28)	200	1												450
D	ATE	TANT	-	¥	a a	,				2 T				4
RUN NO.	OPERATOR DSE  HEATING RATE 15	ATM. 72 0 STANT			·									0
RUN	OPER									D.				400
		mg.												0 PLES)
10		$\mathcal{C}$									NI a			00 250 300 350 FOR CHROMEL ALUMEL THERMOCOUPLES)
Y-AXIS	SCALE # mg. inch (SCALE SETTING X 2)	NOIS												HERM
	SCALE SETTIN	SUPPRESSION.					¥							300 MEL TH
	SCA	SUP								7	4		.0	O
	o inch				c									250 ROMEL
	4			5080	2 0930							t		O CH
X-AXIS	TEMP. SCALE SU	TIME SCALE (ALT.)		J. <b>J</b> .	0830							2. 1		N
	SCA	SCAL										I-	<b>1</b>	RRECT
	TEMP.	TIME												150 T. °C (CORRECTED
														1
	2													100
	Resir	mg.		,							,	,		
 E.	Close #4	1												20
SAMPI	Close #	SIZEZI.6		2540-	,			29						
		,						.gm ,TF	MEICH					0

	SAM	SAMPLE:						X-AXIS	S				Y-AXIS	IS		RUN	RUN NO.	٥	DATE	
	0	#3	<u></u>			- E	TEMP. SCALE	ALE.	Ĭ	o°	SC	SCALE	Ε	D		OPE	OPERATOR	2		
	3					SHIFT	F		inch	inch	(\$C'	(SCALE SETTING X 2)	in FING X 2)	Ę.		HEAT!	HEATING RATE ATM.	ATE		O E
	SIZE	SIZE 22, 3 mg.	3 mg.			IM	E SCAI	TIME SCALE (ALT.)	(.)		SU	SUPPRESSION	NOIS		mg.		TIME CONSTANT	STANT		sec
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	7/							Shel		*					27					
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0		20	0	5	100 T. °C	150 C (COR	150 Z		FOR CHROMEL	250 ROMEL		300 350	OC THERM	3	350	4(	400	4	450	

	SAMI	PLE:					X-AXIS					Y-AXIS	S		RUN NO.	NO.	Q	DATE 7/29	124
	S €	# 3	Dow #6		TEM	TEMP. SCALE 50	LE ST		0 400	SCA	SCALE 4	mg.			OPE HEAT	OPERATOR DE HEATING RATE	TSE /S	\\\\	0.
	. د	1			SHIFT	0	0	÷.	į	(SCA	(SCALE SETTING X 2)	NG X 2)	1	,	ATM.	ATM. NZ (0)	(0)	100	E .
	SIZE	SIZE 71 7 mg.	mg.		Z.	TIME SCALE (ALT.)	E (ALT	0		SUF	SUPPRESSION	NOIS	0	mg.		TIME CONSTANT	TANT	, _	sec
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		50		100 T.	150 T. °C (CORRECTED	150 CORREC	10	00 25C FOR CHROMEL	250 IROMEL	10	300 350 ALUMEL THERMOCOUPLES)	O THERM	35	350 DUPLES)	4(	400	4	450	

19/7			Se													)6 A	
DATE 7/9/7	3 4	?	1 77 L		· · · · ·	-		-		-		-					450
	7	KAIE.	STAN.									77					
RUN NO.	OPERATOR TEL	HEALING KALE TO	AIM. N.2. & 50 cc/ mb								- "					• • • •	00
RUN	OPE	HEX		-11	٠.									*			400
	6 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		mg.	,										,			0
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Y-AXIS	m	incl NG X 2)	NOIS														0
	LE. Y	LE SETTI	SUPPRESSION												l l		300
	SCA	(SCAI	SUP			Fart			b) on			·					
	00	inch	1			Hold			- 1 hr hold					3			250
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X-AXIS	E 50	.=	(ALT	ŀ								***					200
×	TEMP. SCALE	5	0						•				, ,				
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× ×	}			-								•1				-	100
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SAMPLE: Comb Enoxy Res.	Story 3th			-													. 50
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13. ABSTRACT			
The material used as a binder in the			
and Mk 45 Aircraft Parachute Flares,	since the 1968 c	conversion	from Laminac to
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epoxy, has been procured on a sole-source basis from the Dow Chemical Company. A thorough analysis of all pertinent properties of an epoxy produced by Union Carbide Corporation has demonstrated that this material is, in all significant respects, as good as that obtained from Dow. This report recommends that Union Carbide Corporation be approved as an alternate supplier for the epoxy used in these flare candles.

DD FORM 1473 (PAGE 1)

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(PAGE 2)